

Europäisches Patentamt
European Patent Office
Office européen des brevets



(11) **EP 0 823 744 A2**

(12) **EUROPEAN PATENT APPLICATION**

(43) Date of publication:
11.02.1998 Bulletin 1998/07

(51) Int Cl.⁶ **H01M 10/40, H01B 1/12**

(21) Application number: 97305965.2

(22) Date of filing: 06.08.1997

(84) Designated Contracting States:
**AT BE CH DE DK ES FI FR GB GR IE IT LI LU MC
NL PT SE**

(30) Priority: 07.08.1996 JP 224534/96

(71) Applicant: **mitsui petrochemical
industries, ltd.**
Tokyo 100 (JP)

(72) Inventors:
• **Taniuchi, Masahiro**
Ohta-ku, Tokyo 143 (JP)
• **Inoue, Tomohiro**
Ohta-ku, Tokyo 143 (JP)

- **Ohsawa, Toshiyuki**
Ohta-ku, Tokyo 143 (JP)
- **Yokoyama, Keiichi,**
Mitsui Petrochemical Ind. Ltd.
Sodegaura-shi, Chiba 299-02 (JP)
- **Hiwara, Akio, Mitsui Petrochemical Ind. Ltd.**
Sodegaura-shi, Chiba 299-02 (JP)
- **Toriida, Masahiro, Mitsui Petrochemical Ind. Ltd.**
Sodegaura-shi, Chiba 299-02 (JP)

(74) Representative: **Cresswell, Thomas Anthony**
J.A. KEMP & CO.
14 South Square
Gray's Inn
London WC1R 5LX (GB)

(54) **Ionically conductive polymeric gel electrolyte and solid battery containing the same**

(57) An ionically conductive polymeric gel electrolyte comprising a polymer matrix, a non-aqueous elec-

trolytic solution and an electrolytic salt, wherein the solvent of the non-aqueous electrolytic solution comprises a halogen-substituted carbonic ester.

EP 0 823 744 A2

DescriptionFIELD OF THE INVENTION

5 The present invention relates to novel gel electrolytes and solid batteries using the gel electrolytes.

BACKGROUND OF THE INVENTION

10 Batteries are each mainly constituted of positive electrode, negative electrode and electrolytic solution. In order to prevent decrease of lifetime of batteries caused by leakage or vaporization of the electrolytic solution, containers of batteries have been desired to have structures of high rigidity, high sealing properties and excellent pressure resistance (e.g., cylindrical type, square type, coin type). Particularly these days, various forms have been required for batteries, and development of flat type batteries having large area has been made.

15 In this connection, solid batteries have been studied. As solid electrolytes contained in the solid batteries, various ceramics, inorganic conductive glasses (e.g., NASIKON, LISICON), and polymeric solid electrolytes composed of a solid solution of a polymer matrix and electrolytic salt have been studied. As for these materials, however, various problems have been pointed out. For example, the inorganic electrolytes show poor stability or they are restricted in the battery system. The polymeric solid electrolytes have low ionic conductivity or low diaphragm strength, though they show good processability. To cope with these problems, there have been disclosed, for example, polymeric solid elec-
 20 trolytes using polysiloxane (U.S. Patent No. 5,123,512) or polyphosphazene (U.S. Patent 4,840,856) as polymer matrix, and polymeric gel electrolytes using polyethylene oxide-crosslinked products (U.S. Patents No. 5,037,712, No. 5,229,225, No. 5,009,970, No. 5,102,752), ethylene oxide copolymers (U.S. Patent No. 4,816,644, Japanese Patent Laid-Open Publication No. 24164/1991), vinyl copolymers (Japanese Patent Laid-Open Publication No. 320781/1995) or epoxy resins (U.S. Patent No. 5,006,431) as polymer matrix. These electrolytes, however, are still insufficient as
 25 polymeric solid electrolytes for batteries, from the viewpoints of ionic conductivity, film strength and stability.

OBJECT OF THE INVENTION

30 The present invention is intended to solve such problems shown in the solid electrolytes where conventional polymers is matrix as described above, and it is an object of the invention to provide an ionically conductive polymeric gel electrolyte for batteries, which has high ionic conductivity and sufficiently high solid strength. It is another object of the invention to provide a solid battery using the ionically conductive polymeric gel electrolyte, which is excellent in characteristics of preventing internal short-circuit even if no diaphragm is used and which has high reliability.

SUMMARY OF THE INVENTION

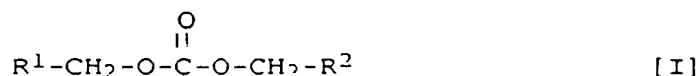
35 The ionically conductive polymeric gel electrolyte of the present invention is characterized in that this gel electrolyte comprises a polymer matrix, a solvent and an electrolytic salt, and that the solvent used for a non-aqueous electrolytic solution contains at least one kind of a halogen-substituted carbonic ester. In general, if a carbonic ester is used as a
 40 solvent of a non-aqueous electrolytic solution of a polymeric gel electrolyte, elastic modulus of the resulting electrolyte for solid battery tends to lower and the retention of the solvent is reduced.

The present inventors have earnestly studied to solve these problems, and as a result, they have found that use of at least one kind of a halogen-substituted carbonic ester as a solvent of the non-aqueous electrolytic solution makes it possible to improve the elastic modulus of the solid electrolyte and to improve the retention of the solvent without
 45 lowering the ionic conductivity. Based on the finding, the present invention has been accomplished.

DETAILED DESCRIPTION OF THE INVENTION

50 The ionically conductive polymeric gel electrolyte of the invention is described in detail hereinafter. The polymeric gel electrolyte of the invention is of a homogeneous viscoelastic body as a whole which comprises a polymer matrix, a solvent and an electrolytic salt. In the present invention, a halogen-substituted carbonic ester having a relative dielectric constant of not less than 4.0 is preferably used as a solvent, because the ionic conductivity of the resulting polymeric gel electrolyte can be enhanced.

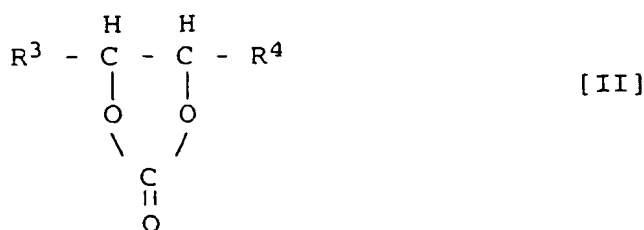
The halogen-substituted non-cyclic carbonic ester for use in the invention is represented by, for example, the
 55 following formula [I]:



wherein R^1 and R^2 are each a halogen-substituted or unsubstituted alkyl group of 1 to 4 carbon atoms, hydrogen or halogen, with the proviso that at least one halogen is present in at least one of R^1 and R^2 ; and R^1 and R^2 may be the same or different.

When R^1 and R^2 in the formula [I] are each an unsubstituted alkyl group of 1 to 4 carbon atoms, the unsubstituted alkyl group is preferably methyl, ethyl or the like. When R^1 and R^2 are each a halogen-substituted alkyl group of 1 to 4 carbon atoms, the halogen-substituted alkyl group is preferably a group which is substituted with halogen at the 1- or more position, such as chloromethyl, 2-chloroethyl, dichloromethyl, 2,2-dichloroethyl, trichloromethyl, 2,2,2-trichloroethyl, pentachloromethyl, 2,2,3,3,3-pentachloropropyl, heptachloromethyl, 2,2,3,3,4,4,4-heptachlorobutyl, fluoromethyl, 2-fluoromethyl, difluoromethyl, 2,2-difluoroethyl, trifluoromethyl, 2,2,2-trifluoroethyl, pentafluoromethyl, 2,2,3,3,3-pentafluoropropyl, bromomethyl, 2-bromoethyl, dibromomethyl, 2,2-dibromoethyl, tribromomethyl, 2,2,2-tribromoethyl, iodomethyl or 2-iodoethyl. When R^1 and R^2 are each a halogen atom, examples of the halogen atoms include fluorine, chlorine, bromine and iodine. Particularly preferable is a fluorine-substituted ester, because cycle properties of the resulting battery can be improved.

The halogen-substituted cyclic carbonic ester for use in the invention is, for example, a carbonic ester represented by the following formula [II]:



wherein R^3 and R^4 are each a halogen-substituted or unsubstituted alkyl group of 1 to 2 carbon atoms, hydrogen or halogen, with the proviso that at least one halogen is present in at least one of R^3 and R^4 ; and R^3 and R^4 may be the same or different.

When R^3 and R^4 in the formula [II] are each an unsubstituted alkyl group of 1 to 2 carbon atoms, the unsubstituted alkyl group is preferably methyl. When R^3 and R^4 are each a halogen-substituted alkyl group of 1 to 2 carbon atoms, the halogen-substituted alkyl group is preferably a group which is substituted with halogen at the 1- or more position, such as chloromethyl, dichloromethyl, trichloromethyl, fluoromethyl, difluoromethyl, trifluoromethyl, bromomethyl, dibromomethyl, tribromomethyl or iodomethyl. When R^1 and R^2 are each a halogen atom, examples of the halogen atoms include fluorine, chlorine, bromine and iodine. Particularly preferable is a fluorine-substituted ester, because cycle properties of the resulting battery can be improved.

Further, use of the halogen-substituted carbonic ester contributes to improvement of flame retardance.

For improving the ionic conductivity, the halogen-substituted non-cyclic carbonic ester is particularly preferably employed.

Examples of the halogen-substituted non-cyclic carbonic esters having a relative dielectric constant of not less than 4.0 include methyl-2-chloroethyl carbonate, methyl-2,2,2-trichloroethyl carbonate, methyl-2,2,2-trifluoroethyl carbonate, di(2,2,2-trifluoroethyl) carbonate and methyl-2,2,3,3,3-pentafluoropropyl carbonate.

The halogen-substituted non-cyclic carbonic ester used in the invention is preferably one having a relative dielectric constant of not less than 6.0, and is more preferably one having a dielectric constant of not less than 7.0.

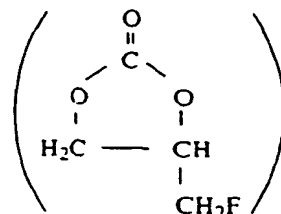
Examples of the halogen-substituted non-cyclic carbonic esters having a relative dielectric constant of not less than 6.0 and less than 7.0 include methyl-2-chloroethyl carbonate and methyl-2,2,3,3,3-pentafluoropropyl carbonate.

Examples of the halogen-substituted non-cyclic carbonic esters having a relative dielectric constant of not less than 7.0 include methyl-2,2,2-trifluoroethyl carbonate and di(2,2,2-trifluoroethyl) carbonate.

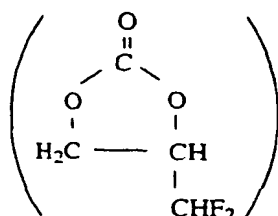
All of the halogen-substituted cyclic carbonic ester of the formula [II] have a relative dielectric constant of not less than 4.0; therefore all of them are preferable.

Examples of the halogen-substituted cyclic carbonic esters include

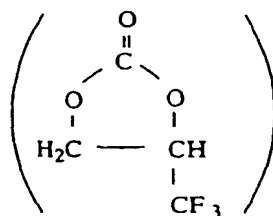
fluoromethylethylene carbonate



difluoromethylethylene carbonate



trifluoromethylethylene carbonate



fluoroethylene carbonate, chloromethylethylene carbonate, dichloromethylethylene carbonate, trichloromethylethylene carbonate, chloroethylene carbonate, bromomethylethylene carbonate, dibromomethylethylene carbonate, tribromomethylethylene carbonate, bromoethylene carbonate, iodomethylethylene carbonate, diiodomethylethylene carbonate, triiodomethylethylene carbonate and iodoethylene carbonate.

In the present invention, the halogen-substituted carbonic esters may be used singly or in combination of two or more kinds.

By the use of the halogen-substituted non-cyclic carbonic ester, a gel electrolyte improved particularly in ionic conductivity, elastic modulus, retention of solvent and low-temperature ionic conductivity can be obtained, and besides a flame-retardant gel electrolyte can be also obtained. In the present invention, substituted cyclic esters can be incorporated as other solvents for forming the electrolytic solution in addition to the halogen-substituted carbonic ester. Examples of such solvents for forming the electrolytic solution include ethylene carbonate (EC) and propylene carbonate (PC). Other solvents include dimethyl carbonate, diethyl carbonate, γ -butylene carbonate, γ -butyrolactone, sulfolane, dioxolane, tetrahydrofuran, 2-methyltetrahydrofuran, methylethyl carbonate, dimethyl sulfoxide, 1,2-dimethoxyethane, 1,2-ethoxymethoxyethane, methyl glyme, methyl triglyme, methyl tetraglyme, ethyl glyme, ethyl diglyme and butyl diglyme. The solvents for forming the electrolytic solution, which are employable in the invention, are not limited to those examples. The above-mentioned solvents may be used singly or in combination.

Of the cyclic esters mentioned above, particularly preferable are cyclic carbonic esters such as EC and PC.

The amount of the halogen-substituted non-cyclic carbonic ester in the mixed solvent of the non-aqueous electrolytic solution is preferably 5 to 80 % by volume, more preferably 20 to 70 % by volume, particularly preferably 30 to 50 % by volume.

The amount of the halogen-substituted cyclic carbonic ester in the mixed solvent of the non-aqueous electrolytic

solution is preferably 20 to 95 % by volume, more preferably 30 to 80 % by volume, particularly preferably 50 to 70 % by volume.

The gel electrolyte of the invention is, for example, a thermoplastic gel composition comprising a polymer (e.g., polyvinylidene fluoride, polyacrylonitrile or polyethylene oxide), a solvent and an electrolytic salt; or a gel composition comprising polysiloxane having an ethylene oxide chain in its side chains or a crosslinked polymer having an ethylene oxide chain in its side chains or main chains (e.g., polymer matrix crosslinked with urethane compounds, acrylic compounds or epoxy compounds), a solvent and an electrolytic salt. Of these, a gel composition having the crosslinked polymer is particularly effective.

The crosslinked polymer matrix is now described in detail. The polymerizable compound for use in the invention contains, in its molecule, a hetero atom other than a carbon atom, such as an oxygen atom, a nitrogen atom or a sulfur atom. The polymeric gel electrolyte is obtained by, for example, dissolving the hetero atom-containing polymerizable compound in a non-aqueous solvent (or electrolytic solution), followed by performing polymerization reaction. The polymerization reaction is conducted preferably in an inert gas atmosphere. There is no specific limitation on the polymerizable compound for use in the invention, and any compounds capable of undergoing polymerization reaction such as thermal polymerization or active light polymerization (photopolymerization) to produce polymers can be employed.

Examples of the polymerizable compounds for use in the invention include monofunctional and polyfunctional (meth)acrylate monomers, and prepolymers thereof. The term "(meth)acrylate" used herein means acrylate or methacrylate.

Examples of the monofunctional acrylates include alkyl (meth)acrylates, such as methyl (meth)acrylate, butyl (meth)acrylate, trifluoroethyl (meth)acrylate, alicyclic (meth)acrylates, hydroxyalkyl (meth)acrylates, such as hydroxyethyl acrylate and hydroxypropyl acrylate; hydroxypolyoxyalkylene (meth)acrylate, such as hydroxypolyoxyethylene (meth)acrylate and hydroxypolyoxypropylene (meth)acrylate; and alkoxyalkyl (meth)acrylates, such as methoxyethyl acrylate, ethoxyethyl acrylate and phenoxyethyl acrylate. In the above hydroxypolyoxyalkylene (meth)acrylates, the oxyalkylene group preferably has 1 to 4 carbon atoms. In the above alkoxyalkyl (meth)acrylates, the alkoxy group preferably has 1 to 4 carbon atoms.

Examples of other (meth)acrylates include ethylene glycol (meth)acrylates, such as methylethylene glycol (meth)acrylate, ethylethylene glycol (meth)acrylate, propylethylene glycol (meth)acrylate, phenylethylene glycol (meth)acrylate, methyl-diethylene glycol (meth)acrylate, ethyl-diethylene glycol (meth)acrylate, methylethoxydiethylene glycol acrylate, methoxydiethylene glycol methacrylate, methoxytriethylene glycol acrylate, methoxytriethylene glycol methacrylate and methoxytetraethylene glycol methacrylate; and propylene glycol (meth)acrylates, such as ethylpropylene glycol acrylate, butylpropylene glycol acrylate and methoxypropylene glycol acrylate.

The (meth)acrylates may contain a heterocyclic group. The heterocyclic group is a residual group of a heterocyclic ring containing a hetero atom such as oxygen, nitrogen or sulfur. There is no specific limitation on the kind of the heterocyclic group contained in the (meth)acrylates, but preferable are (meth)acrylates having a furfuryl group or a tetrahydrofurfuryl group, such as furfuryl (meth)acrylate or tetrahydrofurfuryl (meth)acrylate. Examples of other (meth)acrylates having a heterocyclic group include alkylene glycol acrylates having a furfuryl group or a tetrahydrofurfuryl group, such as furfurylethylene glycol (meth)acrylate, tetrahydrofurfurylethylene glycol (meth)acrylate, furfurylpropylene glycol (meth)acrylate and tetrahydrofurfurylpropylene glycol (meth)acrylate.

The (meth)acrylate compounds or the prepolymers thereof have a molecular weight of usually not more than 1,000, preferably not more than 500, more preferably not more than 300.

In case of a polymeric gel electrolyte obtained by the use of a (meth)acrylate monomer having a molecular weight of more than 1,000, the non-aqueous solvent easily oozes out. The (meth)acrylate compounds mentioned above may be used alone or in combination of two or more kinds.

The (meth)acrylate compound is used in an amount of usually not more than 50 % by weight, preferably 5 to 40 % by weight, more preferably 10 to 30 % by weight, based on the amount of the non-aqueous electrolytic solution.

The polyfunctional (meth)acrylate compound is, for example, a monomer having two or more (meth)acryloyl groups or a prepolymer thereof. Examples of such polyfunctional (meth)acrylate compounds include ethylene glycol dimethacrylate, diethylene glycol di(meth)acrylate, tetraethylene glycol di(meth)acrylate, triethylene glycol di(meth)acrylate, tripropylene glycol diacrylate, EO-modified trimethylolpropane triacrylate, PO-modified trimethylolpropane triacrylate, butanediol (meth)acrylate, trimethylolpropane tri(meth)acrylate and dipentaerythritol hexa(meth)acrylate.

In the preparation of the crosslinked polymer matrix, a combination of a monofunctional monomer and a polyfunctional monomer is particularly preferably used.

If the monofunctional monomer and the polyfunctional monomer are used in combination and if the polyfunctional monomer is a polyfunctional (meth)acrylate compound, the polyfunctional (meth)acrylate compound is used in an amount of not more than 4 % by weight, preferably 0.05 to 2 % by weight, based on the amount of the non-aqueous electrolytic solution.

In the polymerization reaction of the above monomers, a polymerization initiator such as a photopolymerization

initiator or a thermal polymerization initiator is generally employed.

Examples of the photopolymerization initiators include carbonyl compounds, such as benzoin (e.g., benzoin, benzoin methyl ether, benzoin ethyl ether, benzoin isopropyl ether, benzoin isobutyl ether, α -phenylbenzoin) and anthraquinones (e.g., anthraquinone, methylanthraquinone, chloroanthraquinone), other carbonyl compounds (e.g., benzoyldiacetyl, acetophenone, benzophenone, methylbenzoyl formate); sulfur compounds, such as diphenyl sulfide and dithiocarbamate; halides of polycondensed cyclic hydrocarbons, such as α -chloromethylnaphthalene; dyes, such as acriflavine and fluorescein; metallic salts, such as iron chloride and silver chloride; and onium salts, such as p-methoxybenzenediazonium, hexafluorophosphate, diphenyliodonium and triphenylsulfonium. These photopolymerization initiators may be used singly or in combination of two or more kinds. Of the photopolymerization initiators, preferable

are carbonyl compounds, sulfur compounds and onium salts.

Examples of the thermal polymerization initiators include azobisisobutyronitrile, azobisisovaleronitrile, benzoyl peroxide, lauroyl peroxide, ethyl methyl ketone peroxide, bis-(4-t-butylcyclohexyl) peroxydicarbonate and diisopropyl peroxydicarbonate. A sensitizing agent and a storage stabilizer can be optionally used in combination with the polymerization initiator. Further, the thermal polymerization initiator and the photopolymerization initiator may be used in combination.

Examples of the sensitizing agents preferably used in the invention include urea, nitrile compounds such as N,N-di-p-aminobenzonitrile, and phosphorus compounds such as tri-n-butylphosphine. Examples of the storage stabilizers preferably used in the invention include quaternary ammonium chloride, benzothiazole and hydroquinone. The polymerization initiator is used in an amount of usually 0.1 to 10 % by weight, preferably 0.5 to 7 % by weight, based on the total amount of the polymerizable compound. The sensitizing agent or the storage stabilizer is used in an amount of usually 0.1 to 5 parts by weight, based on 100 parts by weight of the polymerizable compound.

Gelation of the electrolyte of the invention can be achieved by pouring into a sealed container the non-aqueous electrolytic solution comprising a polymerizable composition containing the acrylate monomer, or by coating a substrate (e.g., film, metal or glass) with the non-aqueous electrolytic solution and performing polymerization reaction by means of heat or active light. Examples of the active lights generally used in the invention include visible light, ultraviolet light, electron rays and X rays. The resulting polymeric gel electrolyte independently is viscoelastic, and preferably has an elastic modulus of not less than 10^2 dyne/cm when it is used to form a battery, while its ionic conductivity is maintained at not less than 10^{-3} S/cm. The polymeric gel electrolyte can effectively function as a polymeric electrolytic layer of a battery. The gelled electrolyte is made to be in the form of a film, a sheet, or a composite of the electrolyte with a part of other members (constituents) for forming a battery, to produce an article.

Examples of the electrolytic salts for use in the invention include Lewis acid complex salts, such as LiBF_4 , LiAsF_6 , LiPF_6 and LiSbF_6 ; and sulfonic acid electrolytic salts, such as LiCF_3SO_3 , $\text{LiN}(\text{CF}_3\text{SO}_2)_2$, $\text{LiC}(\text{CF}_3\text{SO}_2)_3$, $\text{LiC}(\text{CH}_3)(\text{CF}_3\text{SO}_2)_2$, $\text{LiCH}(\text{CF}_3\text{SO}_2)_2$, $\text{LiCH}_2(\text{CF}_3\text{SO}_2)$, $\text{LiC}_2\text{F}_5\text{SO}_3$, $\text{LiN}(\text{C}_2\text{F}_5\text{SO}_2)_2$, $\text{LiB}(\text{CF}_3\text{SO}_2)_2$ and $\text{LiO}(\text{CF}_3\text{SO}_2)$. There is no limitation to those examples. Other electrolytic salt includes LiClO_4 , LiCF_3CO_3 , NaClO_3 , NaBF_4 , NaSCN , KBF_4 , $\text{Mg}(\text{ClO}_4)_2$ and $\text{Mg}(\text{BF}_4)_2$. The electrolytic salts employable in the invention are not limited to those examples either, and any electrolytic salts used for the conventional non-aqueous electrolytic solutions can be employed. The electrolytic salts mentioned above may be used in combination.

The non-aqueous electrolytic solution is a solution of the electrolytic salt in the aforesaid non-aqueous solvent. The concentration of the electrolytic salt in the non-aqueous electrolytic solution is in the range of usually 1.0 to 7.0 mol per liter of the non-aqueous solvent, preferably 1.0 to 5.0 mol per liter of the non-aqueous solvent. The amount of the non-aqueous electrolytic solution is usually not less than 200 % by weight, preferably 400 to 900 % by weight, particularly preferably 500 to 800 % by weight, based on the amount of the high-molecular weight compound (polymer) for forming the matrix. If the amount thereof is less than 200 % by weight, the resulting electrolyte does not have a sufficiently high ionic conductivity. If the amount thereof exceeds 900 % by weight, solidification of the non-aqueous electrolytic solution becomes difficult.

The polymeric gel electrolyte can be used for polymeric gel electrolytic layers of electrochemical elements, such as batteries, condensers, sensors, electrochromic devices and semiconductor devices.

Next, use of the polymeric gel electrolyte of the invention as a battery electrolyte is described in detail.

In general, a battery consists of a positive electrode made of a positive electrode active material, a negative electrode made of a negative electrode active material, and an electrolyte. By the use of the gel electrolyte of the invention as the battery electrolyte, batteries which have advantageous features unlike the conventional ones can be obtained. When the gel electrolyte of the invention is applied to batteries, the gel electrolyte itself can be made to have also a function of diaphragm. However, the gel electrolyte of the invention is desired to be united to diaphragm into one body for the purpose of producing a uniform electric field between the electrodes, improving strength of the electrolyte and enhancing reliability of the resulting batteries. This should be taken into consideration especially in secondary batteries.

Examples of the positive electrode active materials used for the battery of the invention include transition metal sulfide, such as TiS_2 , MoS_2 and Co_2S_5 ; transition metal oxides, such as V_2O_5 , MnO_2 and CoO_2 ; transition metal chalcogen compounds; and complex compounds of these metal compounds and Li (i.e., Li complex oxides), such as

LiMnO₂, LiMn₂O₄, LiCoO₂, LiNiO₂, LiCo_xNi_{1-x}O₂ (0 < x < 1), LiMn_{2-a}X_aO₄ and LiMn_{2-a-b}X_aY_bO₄ (0 < a < 2, 0 < b < 2, 0 < a+b < 2). Examples of electroconductive materials include one-dimensional graphitization products (thermal polymerization products of organic materials); fluorocarbons; graphites; and electroconductive polymers having an electrical conductivity of not less than 10⁻² S/cm, such as polyaniline, polypyrrole, polyazulene, polyphenylene, polyacetylene, polyphthalocyanine, poly-3-methylthiophene, polypyridine and polydiphenylbenzidine, and derivatives of these conductive polymers.

Examples of the negative electrode active materials used for the battery of the invention include metallic materials, such as lithium, lithium-aluminum alloy, lithium-tin alloy and lithium-magnesium alloy; carbons (including graphite type and non-graphite type); carbon-boron substituted substances (BC₂N); and intercalation materials capable of occluding lithium ion, such as tin oxide. Particular examples of the carbons include calcined graphites, calcined pitch, calcined coke, calcined synthetic polymers and calcined natural polymers. In the present invention, the following carbons are employed: (1) insulating or semiconductor carbons obtained by calcining synthetic polymers such as phenol resin and polyimide or natural polymers in a reducing atmosphere at 400 to 800 °C; (2) electroconductive carbons obtained by calcining coal, pitch, synthetic polymers or natural polymers in a reducing atmosphere at 800 to 1,300 °C; and (3) graphite type carbons, such as those obtained by calcining coke, pitch, synthetic polymers or natural polymers in a reducing atmosphere at not lower than 2,000 °C, and natural graphites.

Examples of positive current collectors for use in the invention include metal sheets, metal foils, metal nets, punching metals, expanded metals, metal plated fibers, metallized wires, and nets or nonwoven fabrics made of metal-containing synthetic fibers. Examples of metals used for these positive current collectors include stainless steel, gold, platinum, nickel, aluminum, molybdenum and titanium. Of these, preferably used are aluminum, stainless steel and titanium from the viewpoints of electrical conductivity, chemical stability, electrochemical stability, economical effects and processability.

Though the polymeric gel electrolyte of the invention can be per se used as diaphragm of batteries, it may be used after a filler is dispersed therein or after it is combined with a porous film (separator) to prepare a composite. Examples of the separators include glass fiber filters; nonwoven fabric filters made of fibers of polymers such as polyester, Teflon, Polyflon, polypropylene and polyethylene; and other nonwoven fabric filters made of mixtures of glass fibers and the above polymeric fibers.

It is preferable that the battery of the present invention is manufactured in the form of one united viscoelastic body from the polymeric gel electrolyte and other battery constituents (members) such as electrodes and diaphragm by a process comprising the steps of impregnating those members with a composition for forming polymeric gel electrolyte and performing polymerization under heating or irradiation with active light. Uniting of the polymeric gel electrolyte to battery members may be carried out member by member. Owing to the polymeric gel electrolyte and the battery members thus united, the electrode reaction at the positive or negative electrode and the ion movement can be smoothly made, whereby internal resistance of the battery can be markedly decreased.

Especially when the polymeric gel electrolyte containing the halogen-substituted carbonic ester is used as a battery constituent, batteries, which have small thickness, are free from short-circuit and leakage of solution, show excellent battery properties and are quite safe, can be obtained because of high elastic modulus of the polymeric gel electrolyte and good retention of the solvent.

EFFECT OF THE INVENTION

Claims 1 to 9 exert such an effect that an ionically conductive polymeric gel electrolyte, which is improved in elastic modulus and retention of solvent without bringing about decrease of ionic conductivity, can be obtained. Also, claims 1 to 9 exert such an effect that a gel solid electrolyte, which shows sufficiently high solid strength and high retention of solvent by virtue of a crosslinked polymer matrix used as the matrix, can be prepared.

Claim 10 exerts such an effect that a solid battery of sufficiently high solid strength, which has a small thickness and is free from solution leakage and short-circuit, can be prepared.

EXAMPLE

The present invention will be further described with reference to the following examples, but it should be construed that the invention is in no way limited to those examples.

The non-aqueous solvents and electrolytic salts used in the following examples are those of battery grade having been sufficiently purified to have water content of not more than 100 ppm and having been subjected to treatment of removing oxygen and nitrogen. In the examples, all of the operations were carried out in an inert gas atmosphere. The ionic conductivity was measured in the following manner using an ionic conductivity measuring cell at 25 °C.

That is, a SUS cylindrical container (inner diameter: 20 mm), an inner surface of which was covered with an insulating tape except the inner bottom surface serving as a counter electrode, was filled with a solid electrolyte. Then, a

SUS column-like material (diameter: 18 mm) serving as a working electrode was fixed by press to the surface of the solid electrolyte, to measure the ionic conductivity.

Example 1

Ionicly conductive polymeric solid electrolyte (I)

In a non-aqueous solvent of propylene carbonate/ethylene carbonate/di(2,2,2-trifluoroethyl) carbonate (4/4/2, by volume) was dissolved $\text{LiN}(\text{CF}_3\text{SO}_2)_2$ in an amount of 2.0 mol per liter of the solvent, to prepare an electrolytic solution. This electrolytic solution of 80 parts by weight was heated to 60 °C, and thereto was added 20 parts by weight of polyacrylonitrile. Then, the mixture thus obtained was cooled to 10 °C to solidify it. Measurement of the ionic conductivity of the obtained solid resulted in 2.7×10^{-3} S/cm. The solid had an elastic modulus of 3.0×10^3 dyne/cm².

Comparative Example 1

Solidification of an electrolytic solution was carried out in the same manner as in Example 1, except that di(2,2,2-trifluoroethyl) carbonate of the non-aqueous solvent was replaced with dipropyl carbonate. The resulting solid had an ionic conductivity of 2.3×10^{-3} S/cm and an elastic modulus of 8.0×10^2 dyne/cm².

Example 2

Ionicly conductive polymeric solid electrolyte (II)

In a non-aqueous solvent of propylene carbonate/ethylene carbonate/methyl-2,2,2-trichloroethyl carbonate (relative dielectric constant: 4.9) (3/3/4, by volume) was dissolved $\text{LiC}(\text{CF}_3\text{SO}_2)_3$ in an amount of 1.8 mol per liter of the solvent, to prepare an electrolytic solution. This electrolytic solution of 85 parts by weight was heated to 60 °C, and thereto was added 15 parts by weight of polyvinylidene fluoride. Then, the mixture thus obtained was cooled to 10 °C to solidify it. The resulting solid had an ionic conductivity of 2.4×10^{-3} S/cm and an elastic modulus of 3.5×10^3 dyne/cm².

Comparative Example 2

Solidification of an electrolytic solution was carried out in the same manner as in Example 2, except that methyl-2,2,2-trichloroethyl carbonate of the non-aqueous solvent was replaced with diisopropyl carbonate (relative dielectric constant: 2.5). The resulting solid had an ionic conductivity of 1.9×10^{-3} S/cm and an elastic modulus of 1.0×10^3 dyne/cm².

Example 3

Ionicly conductive polymeric solid electrolyte (III)

In a non-aqueous solvent of ethylene carbonate/methyl-2,2,2-trifluoroethyl carbonate (1/1, by volume) was dissolved LiPF_6 in an amount of 2.0 mol per liter of the solvent, to prepare an electrolytic solution. In this electrolytic solution of 30 parts by weight were dissolved 20 parts by weight of methyldiethylene glycol methacrylate as a monofunctional monomer and 0.06 part by weight of benzoin isobutyl ether as a photopolymerization initiator, to prepare a photopolymerizable solution. The solution was introduced into the ionic conductivity measuring cell and irradiated with ultraviolet light by means of a highpressure mercury vapor lamp to solidify the electrolytic solution. The resulting solid had an ionic conductivity of 3.0×10^{-3} S/cm and an elastic modulus of 4.1×10^3 dyne/cm².

Comparative Example 3

Solidification of an electrolytic solution was carried out in the same manner as in Example 3, except that methyl-2,2,2-trifluoroethyl carbonate of the non-aqueous solvent was replaced with dimethoxyethane. The resulting solid had an ionic conductivity of 9.7×10^{-4} S/cm and an elastic modulus of 6.0×10^2 dyne/cm².

Example 4Ionicly conductive polymeric solid electrolyte (IV)

In a non-aqueous solvent of propylene carbonate/ethylene carbonate/methyl-2-chloroethyl carbonate (2/3/5, by volume) was dissolved LiCF_3SO_3 in an amount of 1.6 mol per liter of the solvent, to prepare an electrolytic solution. In this electrolytic solution of 75 parts by weight were dissolved 15 parts by weight of diethylene glycol dimethacrylate and 0.05 part by weight of benzoin isopropyl ether as a photopolymerization initiator, to prepare a photopolymerizable solution. The solution was solidified in the same manner as in Example 3. The resulting solid had an ionic conductivity of 2.4×10^{-3} S/cm and an elastic modulus of 3.3×10^3 dyne/cm².

Comparative Example 4

Solidification of an electrolytic solution was carried out in the same manner as in Example 4, except that methyl-2-chloroethyl carbonate of the non-aqueous solvent was replaced with dimethoxyethane. The resulting solid had an ionic conductivity of 2.2×10^{-3} S/cm and an elastic modulus of 7.0×10^2 dyne/cm².

Example 5Ionicly conductive polymeric solid electrolyte (V)

In a non-aqueous solvent of propylene carbonate/ethylene carbonate/methyl-2,2,3,3,3-pentafluoropropyl carbonate (2/5/3, by volume) was dissolved $\text{LiN}(\text{CF}_3\text{SO}_2)_2$ in an amount of 2.0 mol per liter of the solvent, to prepare an electrolytic solution. In this electrolytic solution of 86 parts by weight were dissolved 13.8 parts by weight of ethyldiethylene glycol methacrylate as a monofunctional monomer, 0.2 part by weight of PO-modified trimethylolpropane triacrylate as a polyfunctional monomer and 0.05 part by weight of methylbenzoyl formate as a photopolymerization initiator, to prepare a photopolymerizable solution. The solution was solidified in the same manner as in Example 3. The resulting solid had an ionic conductivity of 2.7×10^{-3} S/cm and an elastic modulus of 3.8×10^3 dyne/cm².

Comparative Example 5

Solidification of an electrolytic solution was carried out in the same manner as in Example 5, except that methyl-2,2,3,3,3-pentafluoropropyl carbonate of the non-aqueous solvent was replaced with dipropyl carbonate. The resulting solid had an ionic conductivity of 1.7×10^{-3} S/cm and an elastic modulus of 5.6×10^2 dyne/cm².

Example 6Ionicly conductive polymeric solid electrolyte (VI)

In a non-aqueous solvent of propylene carbonate/ethylene carbonate/methyl-2,2,2-trifluoroethyl carbonate (3/3/4, by volume) was dissolved LiBF_4 in an amount of 1.8 mol per liter of the solvent, to prepare an electrolytic solution. In this electrolytic solution of 85 parts by weight were dissolved 14.9 parts by weight of methoxytriethylene glycol acrylate as a monofunctional monomer, 0.1 part by weight of dipentaerythritol hexamethacrylate as a polyfunctional monomer and 0.05 part by weight of methylbenzoyl formate as a photopolymerization initiator, to prepare a photopolymerizable solution. The solution was solidified in the same manner as in Example 3. The resulting solid had an ionic conductivity of 2.6×10^{-3} S/cm and an elastic modulus of 3.5×10^3 dyne/cm².

Comparative Example 6

Solidification of an electrolytic solution was carried out in the same manner as in Example 6, except that methyl-2,2,2-trifluoroethyl carbonate of the non-aqueous solvent was replaced with dimethoxyethane. The resulting solid had an ionic conductivity of 2.3×10^{-3} S/cm and an elastic modulus of 8.4×10^2 dyne/cm².

Example 7Ionicly conductive polymeric solid electrolyte (VII)

In a non-aqueous solvent of propylene carbonate/ethylene carbonate/methyl-2-chloroethyl carbonate (2/5/3, by

volume) was dissolved LiAsF_6 in an amount of 2.0 mol per liter of the solvent, to prepare an electrolytic solution. In this electrolytic solution of 80 parts by weight were dissolved 14.7 parts by weight of methoxydiethylene glycol acrylate as a monofunctional monomer, 0.3 part by weight of diethylene glycol diacrylate as a polyfunctional monomer and 0.05 part by weight of methylbenzoyl formate as a photopolymerization initiator were dissolved. to prepare a photopolym-
 5 erizable solution. The solution was solidified in the same manner as in Example 3. The resulting solid had an ionic conductivity of 3.2×10^{-3} S/cm and an elastic modulus of 3.1×10^3 dyne/cm².

Example 8

Ionically conductive polymeric solid electrolyte (VIII)

In a non-aqueous solvent of propylene carbonate/ethylene carbonate/di(2,2,2-trifluoroethyl) carbonate (2/3/5, by volume) were dissolved $\text{LiN}(\text{CF}_3\text{SO}_2)_2$ in an amount of 1.8 mol per liter of the solvent and LiPF_6 in an amount of 0.2 mol per liter of the solvent, to prepare an equivalent mixed electrolytic solution. In this electrolytic solution of 85 parts
 15 by weight were dissolved 14.0 parts by weight of methyldiethylene glycol acrylate as a monofunctional monomer, 1.0 part by weight of trimethylolpropane triacrylate as a polyfunctional monomer and 0.05 part by weight of bis(4-t-butyl-cyclohexyl) peroxydicarbonate as a thermal polymerization initiator, to prepare a polymerizable solution. The solution was subjected to polymerization reaction at 50 °C to solidify the electrolytic solution. The resulting solid had an ionic conductivity of 3.4×10^{-3} S/cm and an elastic modulus of 3.2×10^3 dyne/cm².

Comparative Example 7

Solidification of an electrolytic solution was carried out in the same manner as in Example 7, except that di(2,2,2-trifluoroethyl) carbonate of the non-aqueous solvent was replaced with diisopropyl carbonate. The resulting solid had an
 25 ionic conductivity of 2.0×10^{-3} S/cm and an elastic modulus of 7.3×10^2 dyne/cm².

Example 9

Ionically conductive polymeric solid electrolyte (IX)

In a non-aqueous solvent of propylene carbonate/ethylene carbonate/methyl-2-chloroethyl carbonate (3/3/4, by volume) were dissolved LiCF_3SO_3 in an amount of 1.6 mol per liter of the solvent and LiBF_4 in an amount of 0.2 mol per liter of the solvent, to prepare an equivalent mixed electrolytic solution. In this electrolytic solution of 75 parts by weight were dissolved 20 parts by weight of ethoxydiethylene glycol methacrylate as a monofunctional monomer, 5
 35 parts by weight of diethylene glycol dimethacrylate as a polyfunctional monomer and 0.05 part by weight of benzoin isopropyl ether as a photopolymerization initiator, to prepare a photopolymerizable solution. The solution was solidified in the same manner as in Example 3. The resulting solid had an ionic conductivity of 2.4×10^{-3} S/cm and an elastic modulus of 2.9×10^3 dyne/cm².

Example 10

Ionically conductive polymeric solid electrolyte (X)

In a non-aqueous solvent of propylene carbonate/di(2,2,2-trifluoroethyl) carbonate (6/4, by volume) were dissolved
 45 $\text{LiN}(\text{CF}_3\text{SO}_2)_2$ in an amount of 1.5 mol per liter of the solvent and LiAsF_6 in an amount of 0.5 mol per liter of the solvent, to prepare an equivalent mixed electrolytic solution. In this electrolytic solution of 85 parts by weight were dissolved 14.0 parts by weight of methyldiethylene glycol acrylate as a monofunctional monomer and 1.0 part by weight of trimethylolpropane triacrylate as a polyfunctional monomer, to prepare a polymerizable solution. The solution was irradiated with electron rays to solidify the solution. The resulting solid had an ionic conductivity of 3.8×10^{-3} S/cm and
 50 an elastic modulus of 3.6×10^3 dyne/cm².

Example 11

Ionically conductive polymeric solid electrolyte (XI)

In a non-aqueous solvent of trifluoromethylethylene carbonate/ethylene carbonate (7/3, by volume) were dissolved
 55 LiPF_6 in an amount of 1.8 mol per liter of the solvent, to prepare an electrolytic solution. In this electrolytic solution of 86 parts by weight were dissolved 13.8 parts by weight of ethyldiethylene glycol acrylate as a monofunctional monomer,

0.2 part by weight of trimethylolpropane triacrylate as a polyfunctional monomer and 0.06 part by weight of benzoin isopropyl ether as a photopolymerization initiator, to prepare a photopolymerizable solution. The solution was solidified in the same manner as in Example 3. The resulting solid had an ionic conductivity of 1.8×10^{-3} S/cm and an elastic modulus of 1.4×10^4 dyne/cm².

Example 12

Ionically conductive polymeric solid electrolyte (XII)

In a non-aqueous solvent of difluoromethylethylene carbonate/methyl-2,2,2-trifluoroethyl carbonate (5/5 by volume) were dissolved LiPF_6 in an amount of 1.5 mol per liter of the solvent, to prepare an electrolytic solution. In this electrolytic solution of 86 parts by weight were dissolved 13.8 parts by weight of ethyldiethylene glycol acrylate as a monofunctional monomer, 0.2 part by weight of trimethylolpropane triacrylate as a polyfunctional monomer and 0.06 part by weight of benzoin isopropyl ether as a photopolymerization initiator, to prepare a photopolymerizable solution. The solution was solidified in the same manner as in Example 3. The resulting solid had an ionic conductivity of 2.8×10^{-3} S/cm and an elastic modulus of 4.0×10^3 dyne/cm².

Example 13

The gel electrolytes obtained in Examples 1 to 12 and Comparative Examples 1 to 7 were allowed to stand in an inert gas atmosphere, to evaluate retention of the solvent in each gel electrolyte. The results are set forth in Tables 1 and 2.

Example 14

Evaluation of negative electrode battery properties

To a solution obtained by dissolving 2 parts by weight of polyvinylidene fluoride in 58 parts by weight of N-methylpyrrolidone was added 40 parts by weight of coke having been calcined at 2,500 °C, and they were mixed by roll milling in an inert atmosphere to prepare a coating dispersion for negative electrode. The dispersion was applied onto a copper foil (thickness: 20 μm) by means of a wire bar in the atmosphere and dried at 100 °C for 15 minutes to prepare an electrode having a film thickness of 60 μm . Then, a charge and discharge test was performed using the above electrode as a negative electrode, a Li plate as a counter electrode and each of the gel electrolytes prepared in Examples 1 to 12 and Comparative Examples 1 to 7. The charge and discharge test was carried out in the following manner. Using a charge and discharge measuring device of HJ-201 B (available from Hokuto Denko K.K.), a constant current of 1.5 mA was applied to the gel electrolyte until a change of the battery voltage became 0 V, then a constant voltage was applied for 3 hours, to charge the battery. After one hour, the battery was discharged at a current of 1.5 mA so that the battery voltage became 0.6 V. These charge and discharge operations were repeated to evaluate cycle properties of the negative electrode battery. The results are set forth in Tables 1 and 2.

Table 1

	Ionic Conductivity (S/cm)	Elastic Modulus (dyne/cm ²)	Retention of Solvent	Battery Energy Density (mAh/cc) (initial period: 30 cycles)	Battery Property (after 300 cycles)
Ex. 1	2.7×10^{-3}	3.0×10^3	BB	260	BB
Ex. 2	2.4×10^{-3}	3.5×10^3	BB	265	BB
Ex. 3	3.0×10^{-3}	4.1×10^3	BB	267	BB
Ex. 4	2.4×10^{-3}	3.8×10^3	AA	273	AA
Ex. 5	2.7×10^{-3}	3.8×10^3	AA	254	AA
Ex. 6	2.5×10^{-3}	3.5×10^3	AA	270	AA
Ex. 7	3.2×10^{-3}	3.1×10^3	BB	265	AA
Ex. 8	3.4×10^{-3}	3.2×10^3	AA	258	AA

Table 1 (continued)

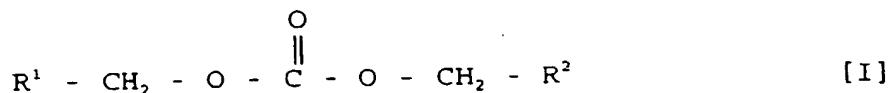
	Ionic Conductivity (S/cm)	Elastic Modulus (dyne/cm ²)	Retention of Solvent	Battery Energy Density (mAh/cc) (initial period: 30 cycles)	Battery Property (after 300 cycles)
Ex. 9	2.4×10^{-3}	2.9×10^3	BB	261	BB
Ex. 10	3.4×10^{-3}	3.2×10^3	AA	270	AA
Ex. 11	1.3×10^{-3}	1.4×10^4	AA	245	BB
Ex. 12	2.3×10^{-3}	4.0×10^3	AA	250	BB
Remarks: "AA" means excellent. "BB" means fairly good. "CC" means good. "DD" means poor.					

Table 2

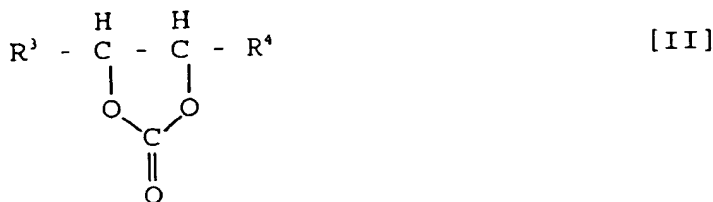
	Ionic Conductivity (S/cm)	Elastic Modulus (dyne/cm ²)	Retention of Solvent	Battery Energy Density (mAh/cc) (initial period: 30 cycles)	Battery Property (after 300 cycles)
Comp. Ex. 1	2.3×10^{-3}	8.0×10^2	DD	260	DD
Comp. Ex. 2	1.9×10^{-3}	1.0×10^3	DD	258	DD
Comp. Ex. 3	9.7×10^{-4}	6.5×10^2	DD	232	CC
Comp. Ex. 4	2.2×10^{-3}	7.0×10^2	DD	270	BB
Comp. Ex. 5	1.7×10^{-3}	5.6×10^2	CC	248	BB
Comp. Ex. 6	2.3×10^{-3}	8.4×10^2	CC	265	AA
Comp. Ex. 7	2.0×10^{-3}	7.3×10^2	CC	250	BB
Remarks: "AA" means excellent "BB" means fairly good. "CC" means good. "DD" means poor.					

Claims

1. An ionically conductive polymeric gel electrolyte comprising a polymer matrix, a non-aqueous electrolytic solution and an electrolytic salt, wherein the solvent of the non-aqueous electrolytic solution comprises a halogen-substituted carbonic ester.
2. An electrolyte as claimed in claim 1, wherein the halogen-substituted carbonic ester is non-cyclic.
3. An electrolyte as claimed in claim 1 or claim 2, wherein the solvent of the non-aqueous electrolytic solution further comprises a cyclic carbonic ester.
4. An electrolyte as claimed in any one of claims 1 to 3, wherein the halogen-substituted carbonic ester has a relative dielectric constant of not less than 4.0.
5. An electrolyte as claimed in any one of claims 1 to 4, wherein the halogen-substituted carbonic ester is a halogen-substituted non-cyclic carbonic ester of formula [I] or a halogen-substituted cyclic carbonic ester of formula [II];



wherein R^1 and R^2 are each a halogen-substituted or unsubstituted alkyl group of 1 to 4 carbon atoms, hydrogen or halogen, with the proviso that at least one halogen is present in at least one of R^1 and R^2 ;



wherein R^3 and R^4 are each a halogen-substituted or unsubstituted alkyl group of 1 to 2 carbon atoms, hydrogen or halogen, with the proviso that at least one halogen is present in at least one of R^3 and R^4 .

6. An electrolyte as claimed in claim 5, wherein the halogen-substituted carbonic ester of formula [I] or formula [II] is a fluorine-substituted carbonic ester.
7. An electrolyte as claimed in any one of claims 1 to 6, wherein the polymer matrix is a crosslinked polymer matrix.
8. An electrolyte as claimed in any one of claims 1 to 7, wherein the non-aqueous electrolytic solution is contained in an amount of at least 200% by weight based on the amount of the polymer matrix.
9. An electrolyte as claimed in any one of claims 1 to 8, wherein the polymer matrix is derived from monomer units which include units of an acrylate monomer modified with an alkylene oxide.
10. A solid battery comprising an electrolyte as claimed in any one of claims 1 to 9.

(19)



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11)

EP 0 823 744 A3

(12)

EUROPEAN PATENT APPLICATION

(88) Date of publication A3:

12.01.2000 Bulletin 2000/02

(51) Int Cl.7: **H01M 10/40, H01B 1/12**

(43) Date of publication A2:

11.02.1998 Bulletin 1998/07

(21) Application number: **97305965.2**

(22) Date of filing: **06.08.1997**

(84) Designated Contracting States:

**AT BE CH DE DK ES FI FR GB GR IE IT LI LU MC
NL PT SE**

(30) Priority: **07.08.1996 JP 22453496**

(71) Applicant: **mitsui chemicals, inc.**
Tokyo (JP)

(72) Inventors:

- **Taniuchi, Masahiro**
Ohta-ku, Tokyo 143 (JP)
- **Inoue, Tomohiro**
Ohta-ku, Tokyo 143 (JP)

- **Ohsawa, Toshiyuki**
Ohta-ku, Tokyo 143 (JP)
- **Yokoyama, Keiichi,**
Mitsui Petrochemical Ind. Ltd.
Sodegaura-shi, Chiba 299-02 (JP)
- **Hiwara, Akio, Mitsui Petrochemical Ind. Ltd.**
Sodegaura-shi, Chiba 299-02 (JP)
- **Toriida, Masahiro, Mitsui Petrochemical Ind. Ltd.**
Sodegaura-shi, Chiba 299-02 (JP)

(74) Representative: **Cresswell, Thomas Anthony**
J.A. KEMP & CO.
14 South Square
Gray's Inn
London WC1R 5LX (GB)

(54) **Ionically conductive polymeric gel electrolyte and solid battery containing the same**

(57) An ionically conductive polymeric gel electrolyte comprising a polymer matrix, a non-aqueous elec-

trolytic solution and an electrolytic salt, wherein the solvent of the non-aqueous electrolytic solution comprises a halogen-substituted carbonic ester.

EP 0 823 744 A3



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 97 30 5965

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.6)
X	US 5 529 859 A (SHU ZHI X ET AL) 25 June 1996 (1996-06-25)	1-6, 10	H01M10/40 H01B1/12
Y	* column 5, line 64 - column 6, line 48 * * column 7, line 7-11; claims 1-14 *	7-9	
Y	US 5 223 353 A (KIMURA OKITOSHI ET AL) 29 June 1993 (1993-06-29) * column 2, line 3 - column 3, line 2; claims 1-13 *	7-9	
X	WO 96 21639 A (STANFORD RES INST INT) 18 July 1996 (1996-07-18) * page 15, line 29 - page 18, line 15 * * page 21, line 1 - page 22, line 16 * * page 25, line 8-19; claims 1,8,16,30,37-39 *	1, 10	
E	WO 97 44842 A (STANFORD RES INST INT) 27 November 1997 (1997-11-27) * page 6, line 9-33 * * page 18, line 12-28 * * page 20, line 10 - page 22, line 21 * * page 45; example 25 *	1-6, 10	TECHNICAL FIELDS SEARCHED (Int.Cl.6)
P, X	DE 197 00 656 A (SANYO CHEMICAL IND LTD) 24 July 1997 (1997-07-24) * claims 2,4-10 *	1,5,6	H01M H01B
A	EP 0 599 534 A (MITSUI PETROCHEMICAL IND ; SONY CORP (JP)) 1 June 1994 (1994-06-01) * page 2, line 34-56 * * page 3, line 43 - page 4, line 30; claims 1-15 *	1,2,5,6	
--- -/-			
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 23 November 1999	Examiner De Vos, L
CATEGORY OF CITED DOCUMENTS		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons A : technological background O : non-written disclosure P : intermediate document & : member of the same patent family corresponding document	
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document			

EPO FORM 1503 (11/92) (P04C01)



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 97 30 5965

DOCUMENTS CONSIDERED TO BE RELEVANT			Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl. 6)
Category	Citation of document with indication, where appropriate, of relevant passages			
A	PATENT ABSTRACTS OF JAPAN vol. 095, no. 009, 31 October 1995 (1995-10-31) & JP 07 165750 A (JAPAN ENERGY CORP), 27 June 1995 (1995-06-27) * abstract *	1		
A	PATENT ABSTRACTS OF JAPAN vol. 1996, no. 01, 31 January 1996 (1996-01-31) & JP 07 240232 A (SONY CORP), 12 September 1995 (1995-09-12) * abstract *	1,5,6		
A	PATENT ABSTRACTS OF JAPAN vol. 1996, no. 06, 28 June 1996 (1996-06-28) & JP 08 045544 A (MITSUI PETROCHEM IND LTD; OTHERS: 01), 16 February 1996 (1996-02-16) * abstract *	1-6		
The present search report has been drawn up for all claims				TECHNICAL FIELDS SEARCHED (Int. Cl. 6)
Place of search THE HAGUE		Date of completion of the search 23 November 1999	Examiner De Vos, L	
CATEGORY OF CITED DOCUMENTS			T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date U : document cited in the application L : document cited for other reasons A : technological background O : non-written disclosure F : intermediate document S : member of the same patent family, corresponding document	

EPO FORM 1503 (3.82) (P-34001)

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 97 30 5965

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

23-11-1999

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
US 5529859	A	25-06-1996	US 5571635 A	05-11-1996
US 5223353	A	29-06-1993	JP 5012913 A	22-01-1993
			DE 4190481 C	28-05-1997
			DE 4190481 T	23-04-1992
			JP 5205515 A	13-08-1993
			WO 9114294 A	19-09-1993
WO 9621639	A	18-07-1996	CA 2210126 A	18-07-1996
			EP 0802898 A	29-10-1997
			JP 10512390 T	24-11-1998
			US 5731104 A	24-03-1998
WO 9744842	A	27-11-1997	US 5830600 A	03-11-1998
			CN 1220029 A	16-06-1999
			EP 0906641 A	07-04-1999
DE 19700656	A	24-07-1997	JP 9251861 A	22-09-1997
			US 5750730 A	12-05-1998
			JP 9286785 A	04-11-1997
EP 0599534	A	01-06-1994	CA 2103298 A	19-05-1994
			DE 69309861 D	22-05-1997
			DE 69309861 T	11-09-1997
			JP 6219992 A	09-08-1994
			US 5633099 A	27-05-1997
			US 5659062 A	19-08-1997
			US 5847188 A	08-12-1998
			JP 7006786 A	10-01-1995
JP 07165750	A	27-06-1995	JP 2936044 B	23-08-1999
JP 07240232	A	12-09-1995	NONE	
JP 08045544	A	16-02-1996	NONE	

EPO FORM P0409

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82